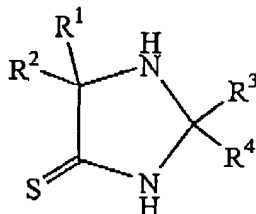


# AMENDMENTS TO THE SPECIFICATION

Page 2, paragraph at lines 6-11:

said method comprising adding to an imidazolidinethione having formula



one of: (i) ~~CHR<sup>5</sup>=CHR<sup>6</sup>-C(Y)ZR<sup>7</sup>~~ CHR<sup>5</sup>=CR<sup>6</sup>-C(Y)ZR<sup>7</sup>; and (ii) R<sup>8</sup>N=C=W to form a reaction mixture; wherein the reaction mixture is substantially free of solvent.

Page 4, paragraph at lines 6-8:

and (b) adding to the imidazolidinethione, without isolation of the imidazolidinethione, one of: (i) ~~CHR<sup>5</sup>=CHR<sup>6</sup>-C(Y)ZR<sup>7</sup>~~ CHR<sup>5</sup>=CR<sup>6</sup>-C(Y)ZR<sup>7</sup>; (ii) R<sup>10</sup>R<sup>11</sup>C=O and R<sup>12</sup>NH<sub>2</sub>; (iii) R<sup>10</sup>R<sup>11</sup>C=NR<sup>12</sup>; and (iv) R<sup>8</sup>N=C=W.

Page 7, paragraph at lines 5-20:

In one embodiment of the invention, the imidazolidinethione reacts with ~~CHR<sup>5</sup>=CHR<sup>6</sup>-C(Y)ZR<sup>7</sup>~~ CHR<sup>5</sup>=CR<sup>6</sup>-C(Y)ZR<sup>7</sup> or R<sup>8</sup>N=C=W substantially in the absence of a solvent. A solvent is any liquid other than the reactants or products of this reaction. Preferably, the reaction mixture contains no more than 5% of solvent by weight, more preferably no more than 2%, more preferably the reaction mixture contains no solvent. Elimination of the solvent increases the efficiency of the process by reducing the cost and the reaction volume. Preferably, the reaction with ~~CHR<sup>5</sup>=CHR<sup>6</sup>-C(Y)ZR<sup>7</sup>~~ CHR<sup>5</sup>=CR<sup>6</sup>-C(Y)ZR<sup>7</sup> or R<sup>8</sup>N=C=W is performed at a temperature from 50°C to 180°C, more preferably from 60°C to 170°C, and most preferably from 90°C to 130°C. The reaction may be followed by well-known methods to determine reaction completion, e.g., IR spectroscopy. Typically, the

reaction is complete in 0.5 to 4 hours. Substitution of acrylate occurs on the thioamide nitrogen or sulfur atom, thereby producing a  $-\text{CHR}^5\text{-CHR}^6\text{-C(Y)ZR}^7$  group as  $\text{B}^1$  or  $\text{B}^2$ , respectively. In contrast, substitution of  $\text{R}^8\text{N=C=W}$  occurs on the amine nitrogen atom of the imidazolidinethione ring, thereby producing a  $-\text{C(W)NHR}^8$  group as  $\text{B}^3$ .

Page 8, paragraph at lines 8-18:

In one embodiment of the invention, an imidazolidinethione is prepared, resulting in a reaction mixture containing the imidazolidinethione, a solvent (typically water or a partially aqueous solvent), and possibly starting materials and byproducts. In this embodiment, one of: (i)  ~~$\text{CHR}^5\text{-CHR}^6\text{-C(Y)ZR}^7$~~   $\text{CHR}^5\text{-CR}^6\text{-C(Y)ZR}^7$ ; (ii)  $\text{R}^{10}\text{R}^{11}\text{C=O}$  and  $\text{R}^{12}\text{NH}_2$ ; (iii)  $\text{R}^{10}\text{R}^{11}\text{C=NR}^{12}$ ; and (iv)  $\text{R}^8\text{N=C=W}$  is added to the reaction mixture without isolation of the imidazolidinethione. Addition of one of these reagents directly to the imidazolidinethione reaction mixture increases the efficiency of the process by eliminating a costly purification step. In one preferred embodiment, the water is partially or substantially completely removed from the reaction mixture prior to addition of one of the aforementioned reagents.